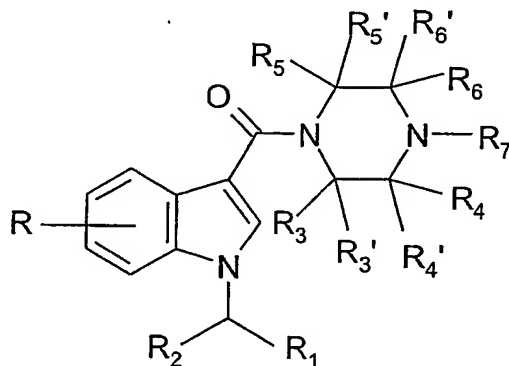


## Claims.

1. An 1-[(indol-3-yl)carbonyl]piperazine derivative having the general formula I



Formula I

wherein

R represents 1-4 substituents independently selected from H, (C<sub>1-4</sub>)alkyl (optionally substituted with halogen), (C<sub>1-4</sub>)alkyloxy (optionally substituted with halogen), halogen, OH, NH<sub>2</sub>, CN and NO<sub>2</sub>;

R<sub>1</sub> is (C<sub>5-8</sub>)cycloalkyl or (C<sub>5-8</sub>)cycloalkenyl;

R<sub>2</sub> is H, methyl or ethyl;

R<sub>3</sub>, R<sub>3'</sub>, R<sub>4</sub>, R<sub>4'</sub>, R<sub>5</sub>, R<sub>5'</sub> and R<sub>6</sub>, R<sub>6'</sub> are independently hydrogen or (C<sub>1-4</sub>)alkyl, optionally substituted with (C<sub>1-4</sub>)alkyloxy, halogen or OH;

R<sub>6</sub> is hydrogen or (C<sub>1-4</sub>)alkyl, optionally substituted with (C<sub>1-4</sub>)alkyloxy, halogen or OH; or

R<sub>6</sub> forms together with R<sub>7</sub> a 4-7 membered saturated heterocyclic ring, optionally containing a further heteroatom selected from O and S;

R<sub>7</sub> forms together with R<sub>6</sub> a 4-7 membered saturated heterocyclic ring, optionally containing a further heteroatom selected from O and S; or

R<sub>7</sub> is H, (C<sub>1-4</sub>)alkyl or (C<sub>3-5</sub>)cycloalkyl, the alkyl groups being optionally substituted with OH, halogen or (C<sub>1-4</sub>)alkyloxy; or

a pharmaceutically acceptable salt thereof.

2. The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 1, wherein R<sub>2</sub> is H and R<sub>1</sub> is (C<sub>5-6</sub>)cycloalkyl.

3. The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 2, wherein R is (C<sub>1-4</sub>)alkyloxy or halogen.

- 4 The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 3, wherein R represents a methoxy group at the 7-position of the indole ring.
- 5 The 1-[(indol-3-yl)carbonyl]piperazine derivative of claim 4, wherein R<sub>3</sub>, R<sub>3</sub>', R<sub>4</sub>', R<sub>5</sub>, R<sub>5</sub>' and R<sub>6</sub>' are H; R<sub>4</sub>, R<sub>6</sub> and R<sub>7</sub> are independently H or (C<sub>1-4</sub>)alkyl; or R<sub>6</sub> forms together with R<sub>7</sub> a 5- or 6-membered saturated heterocyclic ring and R<sub>4</sub> is H or (C<sub>1-4</sub>)alkyl.
6. The 1-[(indol-3-yl)carbonyl]piperazine derivative according to formula I of claim 1 which is selected from:
- 1-[[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl]-3,5-dimethyl-4-ethylpiperazine;
  - 1-[[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl]-3,4,5-trimethylpiperazine;
  - 15 - (S)-1-[[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl]-3,4-dimethylpiperazine;
  - (S)-2-[[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl]-octahydro-2*H*-pyrido-[1, 2-*a*]pyrazine;
  - (S)-2-[[1-(cyclohexylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl]-octahydro-2*H*-pyrrolo-[1, 2-*a*]pyrazine; and
  - 20 - (S)-2-[[1-(cyclopentylmethyl)-7-methoxy-1*H*-indol-3-yl]carbonyl]-octahydro-2*H*-pyrido-[1, 2-*a*]pyrazine;
- or a pharmaceutically acceptable salt thereof.
- 25 7. The 1-[(indol-3-yl)carbonyl]piperazine derivative of any one of claims 1-6 for use in therapy.
8. A pharmaceutical composition comprising an 1-[(indol-3-yl)carbonyl]piperazine derivative of any one of claims 1-6 together with a pharmaceutically acceptable carrier therefor.
- 30 9. Use of an 1-[(indol-3-yl)carbonyl]piperazine derivative of formula I as defined in claim 1, in the preparation of a medicament for the treatment of pain.